

Video AutoFluorescence Imaging (AFI) for dysplasia and cancer in patients with Longstanding Ulcerative Colitis (UC)

Submission date 09/01/2006	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 09/01/2006	Overall study status Completed	☐ Statistical analysis plan ☒ Results
Last Edited 18/07/2008	Condition category Digestive System	☐ Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
NTR498

Study information

Scientific Title

Acronym

AFILUC Study

Study objectives

The aims of the study are:

1. To assess the clinical utility and feasibility of autofluorescence imaging (AFI) in surveillance colonoscopy in patients with longstanding ulcerative colitis (UC)
2. To determine the additional value of AFI in the detection of dysplasia and cancer in these patients
3. To characterise the surface patterns in normal and neoplastic areas in these patients by using narrow band imaging (NBI)

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the local medical ethics committee

Primary study design

Interventional

Study design

Randomised controlled parallel group trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Ulcerative colitis

Interventions

Patients undergoing surveillance colonoscopy with endoscopic tri-modal imaging (ETMI) received inspections of their colonic segments using:

1. Autofluorescence imaging (AFI)
2. White light endoscopy (WLE)

Each patient received both inspections in a random order.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

The value of AFI in patients with UC for detection of dysplasia or cancer.

Key secondary outcome(s)

No secondary outcome measures

Completion date

01/12/2006

Eligibility

Key inclusion criteria

1. Objective diagnosis of UC
2. History of pancolitis
3. Inactive disease determined by a Disease Activity Index

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Known history of colorectal cancer
2. Severe coagulopathy
3. Age less than 18 years

Date of first enrolment

13/04/2005

Date of final enrolment

01/12/2006

Locations

Countries of recruitment

Netherlands

Study participating centre

Academic Medical Center

Amsterdam

Netherlands

1100 DD

Sponsor information

Organisation

Academic Medical Centre (AMC) (The Netherlands)

ROR

<https://ror.org/03t4gr691>

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

Academic Medical Centre (AMC) (The Netherlands)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	01/08/2008		Yes	No